

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification <sup>6</sup> : <b>C07D 239/42, 319/18, 405/12, 409/12, 239/14, A61K 31/505</b>		A1	(11) International Publication Number: <b>WO 95/05366</b>
			(43) International Publication Date: 23 February 1995 (23.02.95)
(21) International Application Number: <b>PCT/EP94/02702</b>		(81) Designated States: AM, AU, BB, BG, BR, BY, CA, CN, CZ, FI, GE, HU, JP, KE, KG, KP, KR, KZ, LK, LT, LV, MD, MG, MN, MW, NO, NZ, PL, RO, RU, SD, SI, SK, TJ, TT, UA, US, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG), ARIPO patent (KE, MW, SD).	
(22) International Filing Date: 12 August 1994 (12.08.94)		Published With international search report.	
(30) Priority Data: 93202445.8 19 August 1993 (19.08.93) EP (34) Countries for which the regional or international application was filed: DE et al. 93202444.1 19 August 1993 (19.08.93) EP (34) Countries for which the regional or international application was filed: DE et al.			
(71) Applicant (for all designated States except US): JANSSEN PHARMACEUTICA N.V. [BE/BE]; Turnhoutseweg 30, B-2340 Beerse (BE).			
(72) Inventors; and (75) Inventors/Applicants (for US only): VAN LOMMEN, Guy, Rosalie, Eugène [BE/BE]; Klets 34, B-2590 Berlaar (BE). DE BRUYN, Marcel, Frans, Leopold [BE/BE]; Pater Schrijverstraat 4, B-2323 Wortel (BE). WIGERINCK, Piet, Tom, Bert, Paul [BE/BE]; Molenstraat 5, B-3010 Leuven (BE).			
(74) Agent: QUAGHEBEUR, Luc; Janssen Pharmaceutica N.V., Patent Dept., Turnhoutseweg 30, B-2340 Beerse (BE).			
(54) Title: VASOCONSTRICTIVE SUBSTITUTED ARYLOXYALKYL DIAMINES			
(57) Abstract			
<p>The present invention is concerned with compounds having formula (I), the pharmaceutically acceptable acid addition salts thereof, and the stereochemically isomeric forms thereof, wherein R<sup>1</sup> is hydrogen or C<sub>1</sub>-alkyl; R<sup>2</sup> is hydrogen or C<sub>1</sub>-alkyl; R<sup>3</sup> is C<sub>1</sub>-alkyl, hydroxy, cyano, halo, C<sub>1</sub>-alkyloxy, aryloxy, arylmethoxy, C<sub>2</sub>-alkenyl, C<sub>2</sub>-alkynyl, C<sub>1</sub>-alkyl-S-, C<sub>1</sub>-alkyl(S=O)-, C<sub>1</sub>-alkylcarbonyl; R<sup>4</sup> is hydrogen, halo, hydroxy, C<sub>1</sub>-alkyl, or C<sub>1</sub>-alkyloxy; or R<sup>3</sup> and R<sup>4</sup> taken together form a bivalent radical; R<sup>5</sup> and R<sup>6</sup> each independently are hydrogen, halo, hydroxy, C<sub>1</sub>-alkyl, C<sub>1</sub>-alkyloxy, aryloxy or arylmethoxy; R<sup>7</sup> is hydrogen; Alk<sup>1</sup> is C<sub>2</sub>-alkanediyl; Alk<sup>2</sup> is C<sub>2</sub>-alkanediyl; Q is a heterocyclic ring containing at least one nitrogen atom or a radical of formula (aa), pharmaceutical compositions, preparations and use as a medicine are described.</p>			
<div style="text-align: center;"> <p>(I)</p> </div> <div style="text-align: center;"> <p>(aa)</p> </div>			

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	GB	United Kingdom	MR	Mauritania
AU	Australia	GE	Georgia	MW	Malawi
BB	Barbados	GN	Guinea	NE	Niger
BE	Belgium	GR	Greece	NL	Netherlands
BF	Burkina Faso	HU	Hungary	NO	Norway
BG	Bulgaria	IE	Ireland	NZ	New Zealand
BJ	Benin	IT	Italy	PL	Poland
BR	Brazil	JP	Japan	PT	Portugal
BY	Belarus	KE	Kenya	RO	Romania
CA	Canada	KG	Kyrgyzstan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SI	Slovenia
CI	Côte d'Ivoire	LI	Liechtenstein	SK	Slovakia
CM	Cameroon	LK	Sri Lanka	SN	Senegal
CN	China	LU	Luxembourg	TD	Chad
CS	Czechoslovakia	LV	Latvia	TG	Togo
CZ	Czech Republic	MC	Monaco	TJ	Tajikistan
DE	Germany	MD	Republic of Moldova	TT	Trinidad and Tobago
DK	Denmark	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	US	United States of America
FI	Finland	MN	Mongolia	UZ	Uzbekistan
FR	France			VN	Viet Nam
GA	Gabon				

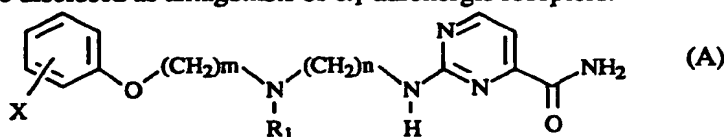
## VASOCONSTRICTIVE SUBSTITUTED ARYLOXYALKYL DIAMINES

5       The present invention relates to novel substituted aryloxyalkyldiamine derivatives, processes for their preparations, pharmaceutical compositions containing them and their use as a medicine, in particular for the prevention and/or treatment of disorders characterized by excessive vasodilatation, especially migraine.

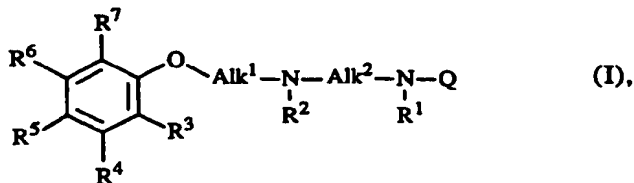
Migraine is a non-lethal disease suffered by one in ten individuals. The main symptom is headache; other symptoms include vomiting and photophobia. For many years the most widely used treatment for migraine involved the administration of ergotalkaloids, which show however several adverse side effects. Recently a tryptamine derivative, i.e. sumatriptan, was introduced as a novel antimigraine drug. We have now surprisingly found that the present novel substituted aryloxyalkyl diamine derivatives show 5-HT<sub>1</sub>-like agonistic activity and can thus be used in the treatment of disorders characterized by excessive vasodilatation, especially migraine.

In Arzneimittel-Forschung, 25, 1404 (1975) some guanidine and amidine derivatives, among which N-[2-[2-(2-methoxyphenoxy)ethylamino]ethyl]guanidine, are disclosed as having noradrenaline depleting activity.

20 In EP-0,511,072 derivatives of 2-aminopyrimidine-4-carboxamide having the general formula (A) are disclosed as antagonists of  $\alpha_1$ -adrenergic receptors.



25      The present invention is concerned with compounds having the formula

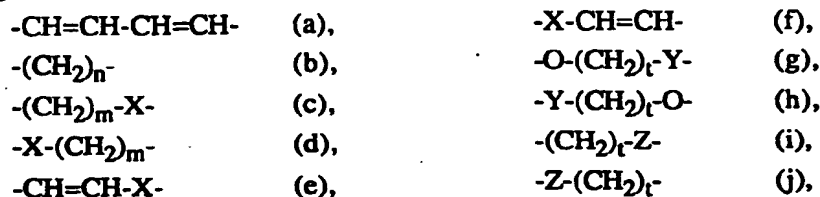


the pharmaceutically acceptable acid addition salts thereof, and the stereochemically isomeric forms thereof, wherein

30 R<sup>1</sup> and R<sup>2</sup> each independently are hydrogen or C<sub>1-6</sub>alkyl;  
R<sup>3</sup> is C<sub>1-6</sub>alkyl, hydroxy, cyano, halo, C<sub>1-6</sub>alkyloxy, aryloxy, arylmethoxy,  
C<sub>2-6</sub>alkenyl, C<sub>2-6</sub>alkynyl, C<sub>1-6</sub>alkyl-S-, C<sub>1-6</sub>alkyl(S=O)-, C<sub>1-6</sub>alkylcarbonyl;

-2-

$R^4$  is hydrogen, halo, hydroxy,  $C_{1-6}$ alkyl, or  $C_{1-6}$ alkyloxy; or  $R^3$  and  $R^4$  taken together form a bivalent radical of formula



in these bivalent radicals one or two hydrogen atoms may be substituted with  $C_{1-6}$ alkyl,  $C_{1-6}$ alkylcarbonyl or  $C_{1-6}$ alkyl-S(O)-;

5 each X independently is -O-, -S-, -S(O)-, -S(O)<sub>2</sub>-, -C(O)-, -NR<sup>8</sup>-;

n is 3 or 4;

each Y independently is -O-, -S-, -S(O)-, -S(O)<sub>2</sub>-, -C(O)-, -NR<sup>8</sup>-;

m is 2 or 3;

each Z is -O-C(O)-, -C(O)-O-, -NH-C(O)-, -C(O)-NH-, -O-S(O)<sub>2</sub>-;

10 t is 1 or 2;

R<sup>8</sup> is hydrogen,  $C_{1-6}$ alkyl,  $C_{1-6}$ alkylcarbonyl or  $C_{1-6}$ alkyl-S(O)-;

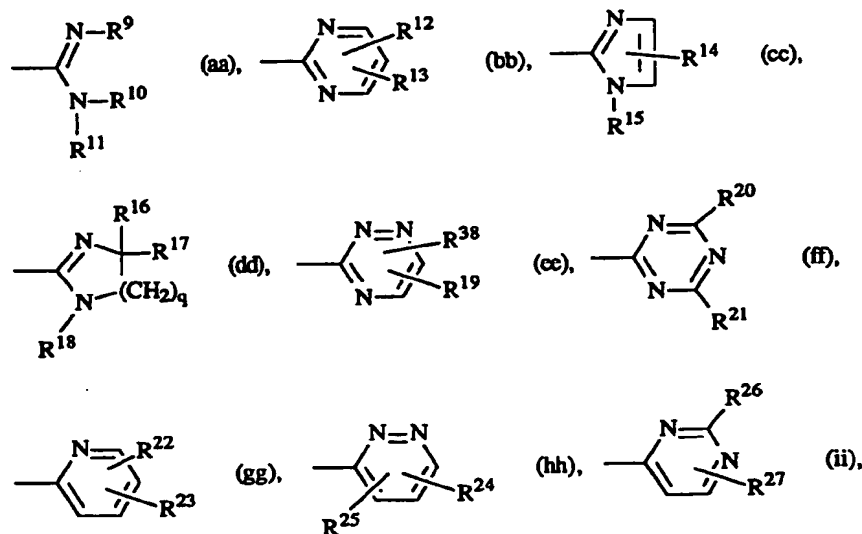
R<sup>5</sup> and R<sup>6</sup> each independently are hydrogen, halo, hydroxy,  $C_{1-6}$ alkyl,  $C_{1-6}$ alkyloxy, aryloxy or arylmethoxy;

R<sup>7</sup> is hydrogen;

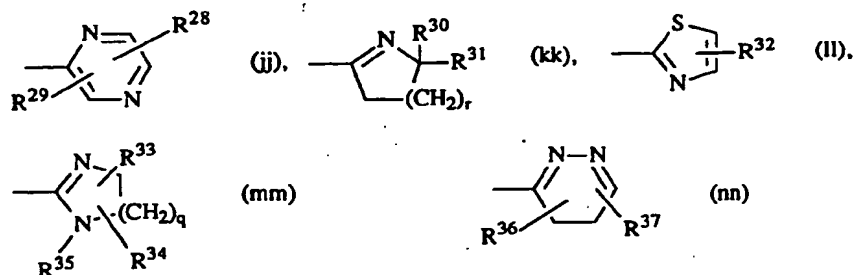
15 Alk<sup>1</sup> is  $C_{2-5}$ alkanediyl;

Alk<sup>2</sup> is  $C_{2-15}$ alkanediyl;

Q is a radical of formula



-3-



5

wherein

R<sup>9</sup> is hydrogen, cyano, aminocarbonyl or C<sub>1-6</sub>alkyl;R<sup>10</sup> is hydrogen, C<sub>1-6</sub>alkyl, C<sub>3-6</sub>alkenyl, C<sub>3-6</sub>alkynyl, C<sub>3-6</sub>cycloalkyl or arylC<sub>1-6</sub>alkyl;R<sup>11</sup> is hydrogen or C<sub>1-6</sub>alkyl; or

10 R<sup>10</sup> and R<sup>11</sup> taken together may form a bivalent radical of formula -(CH<sub>2</sub>)<sub>4</sub>- or -(CH<sub>2</sub>)<sub>5</sub>- , or a piperazine which is optionally substituted with C<sub>1-6</sub>alkyl;

R<sup>12</sup>, R<sup>13</sup>, R<sup>14</sup>, R<sup>19</sup>, R<sup>20</sup>, R<sup>21</sup>, R<sup>22</sup>, R<sup>23</sup>, R<sup>24</sup>, R<sup>25</sup>, R<sup>26</sup>, R<sup>27</sup>, R<sup>28</sup>, R<sup>29</sup>, R<sup>36</sup>, R<sup>37</sup> and R<sup>38</sup> each independently are hydrogen, hydroxy, halo, C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkyloxy, aryloxy, C<sub>1-6</sub>alkylthio, cyano, amino, mono- or di(C<sub>1-6</sub>alkyl)amino, mono- or di(C<sub>3-6</sub>cycloalkyl)-

15 amino, aminocarbonyl, C<sub>1-6</sub>alkyloxycarbonylamino, C<sub>1-6</sub>alkylaminocarbonylamino, piperidinyl, pyrrolidinyl;

R<sup>15</sup>, R<sup>18</sup> and R<sup>35</sup> each independently are hydrogen, C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkylcarbonyl, or arylC<sub>1-6</sub>alkyl;

q is 1, 2 or 3;

20 R<sup>16</sup> and R<sup>17</sup> are both hydrogen, or taken together with the carbon atom to which they are connected form C(O);

r is 1, 2 or 3;

R<sup>30</sup> and R<sup>31</sup> are both hydrogen or taken together with the carbon atom to which they are connected form C(O);

25 R<sup>32</sup> is hydrogen, halo or C<sub>1-6</sub>alkyl;

R<sup>33</sup> is hydrogen and R<sup>34</sup> is hydroxy; or R<sup>33</sup> and R<sup>34</sup> taken together may form a bivalent radical of formula (CH<sub>2</sub>)<sub>3</sub> or (CH<sub>2</sub>)<sub>4</sub> which is optionally substituted with C<sub>1-6</sub>alkyl; and aryl is phenyl optionally substituted hydroxy, halo, C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkyloxy.

30

All the compounds of formula (I) are deemed novel except for

(a) N-[2-[2-(2-methoxyphenoxy)ethylamino]ethyl]guanidine; and

(b) the compounds of formula (I) wherein R<sup>3</sup> is methoxy, ethoxy or isopropyl; R<sup>4</sup> is hydrogen; R<sup>5</sup> is hydrogen; R<sup>6</sup> is chloro, fluoro or methyl; R<sup>7</sup> is hydrogen; R<sup>2</sup> is

-4-

hydrogen or methyl;  $R^1$  is hydrogen;  $Alk^1$  is 1,2-ethanediyl or 1,3-propanediyl;  $Alk^2$  is 1,2-ethanediyl or 1,3-propanediyl; Q is a radical of formula (bb), wherein  $R^{12}$  is hydrogen and  $R^{13}$  is 4-aminocarbonyl.

- 5        Some of the compounds of formula (I) may also exist in their tautomeric forms. Such forms although not explicitly indicated in the above formula are intended to be included within the scope of the present invention.

- As used in the foregoing definitions halo defines fluoro, chloro, bromo and iodo;
- 10         $C_{1-6}alkyl$  defines straight and branch chained saturated hydrocarbon radicals having from 1 to 6 carbon atoms such as, for example, methyl, ethyl, propyl, butyl, pentyl, hexyl as well as the branched isomers thereof;  $C_{3-6}alkenyl$  defines straight and branch chained hydrocarbon radicals containing one double bond and having from 3 to 6 carbon atoms such as, for example, 2-propenyl, 3-butenyl, 2-butenyl, 2-pentenyl, 3-pentenyl,
- 15        3-methyl-2-butenyl and the like; and the carbon atom of said  $C_{3-6}alkenyl$  being connected to a nitrogen atom preferably is saturated;  $C_{2-6}alkenyl$  defines  $C_{3-6}alkenyl$  and the lower homologue thereof, i.e. ethenyl;  $C_{3-6}alkynyl$  defines straight and branch chained hydrocarbon radicals containing one triple bond and having from 3 to 6 carbon atoms such as, for example, 2-propynyl, 3-butyne, 2-butyne, 2-pentyne, 3-pentyne,
- 20        3-hexynyl, and the like; and the carbon atom of said  $C_{3-6}alkynyl$  radical being connected to a nitrogen atom preferably is saturated;  $C_{2-6}alkynyl$  defines  $C_{3-6}alkynyl$  and the lower homologue thereof, i.e. ethynyl;  $C_{3-6}cycloalkyl$  is generic to cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl;  $C_{2-5}alkanediyl$  defines bivalent straight and branch chained saturated hydrocarbon radicals having from 2 to 5 carbon atoms such as, for example,
- 25        1,2-ethanediyl, 1,3-propanediyl, 1,4-butanediyl, 1,5-pentanediyl and the like;  $C_{2-15}alkanediyl$  defines bivalent straight and branch chained saturated hydrocarbon radicals having from 2 to 15 carbon atoms such as, for example, 1,2-ethanediyl, 1,3-propanediyl, 1,4-butanediyl, 1,5-pentanediyl, 1,6-hexanediyl, 1,7-heptanediyl, 1,8-octanediyl, 1,9-nonanediyl, 1,10-decanediyl, 1,11-undecanediyl, 1,12-dodecanediyl,
- 30        1,13-tridecanediyl, 1,14-tetradecanediyl, 1,15-pentadecanediyl, and the branched isomers thereof. The term "C(O)" refers to a carbonyl group.

- The pharmaceutically acceptable acid addition salts as mentioned hereinabove are meant to comprise the therapeutically active non-toxic acid addition salt forms which the
- 35        compounds of formula (I) are able to form. The latter can conveniently be obtained by treating the base form with such appropriate acids as inorganic acids, for example, hydrohalic acids, e.g. hydrochloric, hydrobromic and the like; sulfuric acid; nitric acid; phosphoric acid and the like; or organic acids, for example, acetic, propanoic, hydroxy-

-5-

acetic, 2-hydroxypropanoic, 2-oxopropanoic, ethanedioic, propanedioic, butanedioic, (Z)-2-butenedioic, (E)-2-butenedioic, 2-hydroxybutanedioic, 2,3-dihydroxybutanedioic, 2-hydroxy-1,2,3-propanetricarboxylic, methanesulfonic, ethanesulfonic, benzene-sulfonic, 4-methylbenzenesulfonic, cyclohexanesulfamic, 2-hydroxybenzoic, 4-amino-2-hydroxybenzoic and the like acids. Conversely the salt form can be converted by treatment with alkali into the free base form.

The term addition salt also comprises the hydrates and solvent addition forms which the compounds of formula (I) are able to form. Examples of such forms are e.g. hydrates, alcoholates and the like.

The term "stereochemically isomeric forms" as used hereinbefore defines all the possible isomeric forms which the compounds of formula (I) may possess. Unless otherwise mentioned or indicated, the chemical designation of compounds denotes the mixture of all possible stereochemically isomeric forms, said mixtures containing all diastereomers and enantiomers of the basic molecular structure. More in particular, stereogenic centers may have the R- or S-configuration; C<sub>2-6</sub>-alkenyl radicals may have the E- or Z-configuration. Stereochemically isomeric forms of the compounds of formula (I) are obviously intended to be embraced within the scope of this invention.

R<sup>1</sup> is suitably hydrogen or methyl, preferably R<sup>1</sup> is hydrogen;  
 R<sup>2</sup> is suitably hydrogen or methyl, preferably R<sup>2</sup> is hydrogen;  
 R<sup>3</sup> is suitably C<sub>1-6</sub>-alkyl, hydroxy, C<sub>1-6</sub>-alkyloxy, arylmethoxy, preferably R<sup>3</sup> is methyl, ethyl, hydroxy, methoxy, ethoxy or phenylmethoxy;  
 R<sup>4</sup> is suitably hydrogen, C<sub>1-6</sub>-alkyl or C<sub>1-6</sub>-alkyloxy, preferably R<sup>4</sup> is hydrogen or methoxy;  
 or when taken together  
 R<sup>3</sup> and R<sup>4</sup> form suitably a bivalent radical for formula (a), (b), (e), (f), (g) or (h);  
 each X is suitably O or S, preferably each X is O;  
 each Y is suitably O or S, preferably each Y is O;  
 each Z is suitably -O-C(O)-, -C(O)-O-;  
 R<sup>8</sup> is suitably hydrogen or C<sub>1-6</sub>-alkyl; preferably  
 R<sup>8</sup> is hydrogen or methyl;  
 R<sup>5</sup> is suitably hydrogen or C<sub>1-6</sub>-alkyl, preferably R<sup>5</sup> is hydrogen or methyl;  
 R<sup>6</sup> is suitably hydrogen or C<sub>1-6</sub>-alkyl, preferably R<sup>6</sup> is hydrogen or methyl;  
 Alk<sup>1</sup> is suitably C<sub>2-3</sub>-alkanediyl, preferably Alk<sup>1</sup> is 1,2-ethanediyl, 1,2-propanediyl or 1,3-propanediyl;  
 Alk<sup>2</sup> is suitably C<sub>2-6</sub>-alkanediyl, preferably Alk<sup>2</sup> is 1,3-propanediyl or 1,4-butanediyl;  
 Q is preferably a radical of formula (aa), (bb) or (dd);

- R<sup>9</sup> is suitably hydrogen, cyano, aminocarbonyl or methyl, preferably R<sup>9</sup> is hydrogen or cyano;
- R<sup>10</sup> is suitably hydrogen or C<sub>1-6</sub>alkyl, preferably R<sup>10</sup> is hydrogen, methyl or ethyl;
- R<sup>11</sup> is suitably hydrogen or C<sub>1-6</sub>alkyl, preferably R<sup>11</sup> is hydrogen or methyl;
- 5 R<sup>12</sup> and R<sup>13</sup> each independently are suitably hydrogen, hydroxy, halo or methyl, preferably both R<sup>12</sup> and R<sup>13</sup> are hydrogen or R<sup>12</sup> is hydrogen and R<sup>13</sup> is hydroxy;
- R<sup>14</sup> is suitably hydrogen or hydroxy, preferably R<sup>14</sup> is hydrogen;
- R<sup>15</sup> is suitably hydrogen or phenylmethyl, preferably R<sup>15</sup> is hydrogen;
- q is preferably 2;
- 10 R<sup>16</sup> and R<sup>17</sup> are both preferably hydrogen;
- R<sup>18</sup> is suitably hydrogen or phenylmethyl, preferably R<sup>18</sup> is hydrogen;
- R<sup>19</sup> is suitably hydrogen, halo or methyl, preferably R<sup>19</sup> is hydrogen or chloro;
- R<sup>20</sup> and R<sup>21</sup> each independently suitably are hydrogen, halo or methyl, preferably R<sup>20</sup> and R<sup>21</sup> are hydrogen or chloro;
- 15 R<sup>22</sup> and R<sup>23</sup> each independently suitably are hydrogen, hydroxy, chloro or methyl, preferably R<sup>22</sup> and R<sup>23</sup> are both hydrogen or R<sup>22</sup> is hydrogen and R<sup>23</sup> is hydroxy;
- R<sup>24</sup> and R<sup>25</sup> each independently suitably are hydrogen, hydroxy, halo or methyl, preferably R<sup>24</sup> and R<sup>25</sup> are both hydrogen or R<sup>24</sup> is hydrogen and R<sup>25</sup> is chloro;
- R<sup>26</sup> and R<sup>27</sup> each independently suitably are hydrogen, halo, C<sub>1-6</sub>alkyloxy,
- 20 C<sub>1-6</sub>alkylthio, amino, mono- or di(C<sub>1-6</sub>alkyl)amino; preferably R<sup>26</sup> is hydrogen, chloro, methylthio or amino and R<sup>27</sup> is hydrogen;
- R<sup>28</sup> and R<sup>29</sup> each independently suitably are hydrogen, halo, C<sub>1-6</sub>alkyl, preferably R<sup>28</sup> and R<sup>29</sup> are hydrogen or chloro;
- r preferably is 2;
- 25 R<sup>30</sup> and R<sup>31</sup> both preferably are hydrogen;
- R<sup>32</sup> is suitably hydrogen or methyl, preferably R<sup>32</sup> is hydrogen; and
- aryl is preferably phenyl.

Special compounds of formula (I) are those compounds of formula (I) wherein

30 wherein R<sup>3</sup> is C<sub>1-6</sub>alkyl, hydroxy, C<sub>1-6</sub>alkyloxy, aryloxy, arylmethoxy, C<sub>2-6</sub>alkenyl, C<sub>2-6</sub>alkynyl; one of R<sup>4</sup>, R<sup>5</sup> and R<sup>6</sup> is hydrogen and the others each independently are hydrogen, halo, hydroxy, C<sub>1-6</sub>alkyl, or C<sub>1-6</sub>alkyloxy, Q is a radical of formula (aa), (bb), (cc), (dd), (ee) wherein R<sup>38</sup> is hydrogen, (ff), (gg), (hh), (ii), (jj), (kk), (ll).

35 Other special compounds of formula (I) are those compounds of formula (I) wherein R<sup>3</sup> and R<sup>4</sup> taken together form a bivalent radical of formula



-7-

-CH=CH-CH=CH-	(a),	-X-CH=CH-	(f),
-(CH <sub>2</sub> ) <sub>n</sub> -	(b),	-O-(CH <sub>2</sub> ) <sub>2</sub> -Y-	(g),
-(CH <sub>2</sub> ) <sub>m</sub> -X-	(c),	-Y-(CH <sub>2</sub> ) <sub>2</sub> -O-	(h),
-X-(CH <sub>2</sub> ) <sub>m</sub> -	(d),	-(CH <sub>2</sub> ) <sub>t</sub> -Z-	(i),
-CH=CH-X-	(e),	-Z-(CH <sub>2</sub> ) <sub>t</sub> -	(j),

in these bivalent radicals one or two hydrogen atoms may be substituted with C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkylcarbonyl or C<sub>1-6</sub>alkylsulfoxyl; and wherein X, Y, Z, m, n, and t are defined as in claim 1; and Q is a radical of formula (aa), (bb), (cc), (dd), (ee) wherein R<sup>38</sup> is hydrogen, (ff), (gg), (hh), (ii), (jj), (kk), (ll).

5

Interesting compounds are those compounds of formula (I), wherein R<sup>1</sup> and R<sup>2</sup> both are hydrogen.

10 An interesting subset of compounds are those compounds of formula (I), wherein R<sup>3</sup> and R<sup>4</sup> taken together do not form a bivalent radical and wherein R<sup>3</sup> is C<sub>1-6</sub>alkyl, hydroxy, C<sub>1-6</sub>alkyloxy or arylmethoxy, especially methyl, hydroxy, methoxy, ethoxy and phenylmethoxy.

15 Further interesting compounds are those compounds of formula (I) wherein R<sup>4</sup> is hydrogen, C<sub>1-6</sub>alkyl or C<sub>1-6</sub>alkyloxy and R<sup>5</sup> is hydrogen or C<sub>1-6</sub>alkyloxy.

Particular compounds are those compounds of formula (I) wherein Q is a radical of formula (aa), (bb) or (dd), especially (bb) or (dd).

20

Particularly interesting compounds are those interesting compounds, wherein Q is a radical of formula of (bb), wherein R<sup>9</sup> and R<sup>10</sup> are hydrogen.

25 Another group of particularly interesting compounds are those interesting compounds wherein Q is a radical of formula (dd), wherein q is 2, R<sup>16</sup> and R<sup>17</sup> are hydrogen and R<sup>18</sup> is hydrogen.

30 Another interesting subset of compounds are those compounds of formula (I), wherein R<sup>3</sup> and R<sup>4</sup> taken together form a bivalent radical of formula (a), (b), (e), (f), (g) or (h);

Particular compounds are those compounds of formula (I) wherein Q is a radical of formula (aa), (bb) or (dd), especially (bb) or (dd).

Particularly interesting compounds are those interesting compounds, wherein Q is a radical of formula (bb), wherein R<sup>12</sup> and R<sup>13</sup> are hydrogen.

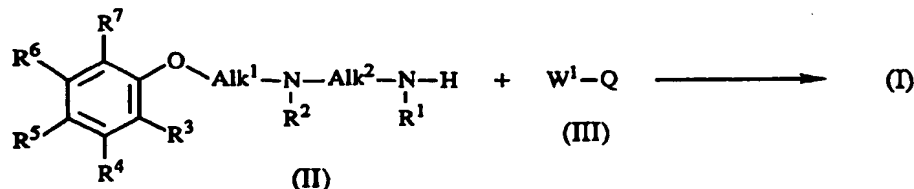
Another group of particularly interesting compounds, wherein Q is a radical of formula (dd), wherein q is 2, R<sup>15</sup> and R<sup>16</sup> are hydrogen and R<sup>17</sup> is hydrogen.

Preferred compounds are :

N-[2-(2,3-dimethoxyphenoxy)ethyl]-N'-2-pyrimidinyl-1,3-propanediamine;  
 2-[2-[[3-(2-pyrimidinylamino)propyl]amino]ethoxy]phenol; N-[2-(2,3-dimethoxy-  
 phenoxy)ethyl]-N'-(1,4,5,6-tetrahydro-2-pyrimidinyl)-1,3-propanediamine;  
 N-[2-(2-methoxyphenoxy)ethyl]-N'-(1,4,5,6-tetrahydro-2-pyrimidinyl)-1,3-propane  
 diamine; N-[2-(2-ethoxyphenoxy)ethyl]-N'-(1,4,5,6-tetrahydro-2-pyrimidinyl)-  
 1,3-propanediamine; N-[3-(2-methoxyphenoxy)propyl]-N'-(1,4,5,6-tetrahydro-2-  
 pyrimidinyl)-1,3-propanediamine; N-[2-[(2,3-dihydro-1,4-benzodioxin-5-yl)oxy]-  
 ethyl]-N'-2-pyrimidinyl-1,3-propanediamine; N-[2-[(2,3-dihydro-1,4-benzodioxin-5-yl)-  
 oxy]ethyl]-N'-(1,4,5,6-tetrahydro-2-pyrimidinyl)-1,3-propanediamine;  
 N-[2-[(2,3-dihydro-1,4-benzodioxin-5-yl)oxy]ethyl]-N'-(1,4,5,6-tetrahydro-2-  
 pyrimidinyl)-1,4-butanediamine; N-[2-(1-naphthalenyloxy)ethyl]-  
 N'-(1,4,5,6-tetrahydro-2-pyrimidinyl)-1,3-propanediamine, the pharmaceutically  
 acceptable acid addition salt thereof or the stereochemically isomeric forms thereof.

The compounds of formula (I) can generally be prepared by reacting a diamine of formula (II) with a reagent of formula (III) wherein W<sup>1</sup> is a reactive leaving group such as, for example, halo, e.g. chloro, bromo; alkyloxy, e.g. methoxy, ethoxy and the like; aryloxy, e.g. phenoxy and the like; alkylthio, e.g. methylthio, ethylthio and the like; arylthio, e.g. benzenethio and the like.

In the formulas (II), (III) and all the following formulas the variables R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, Alk<sup>1</sup>, Alk<sup>2</sup>, and Q are as defined under formula (I), unless indicated otherwise.

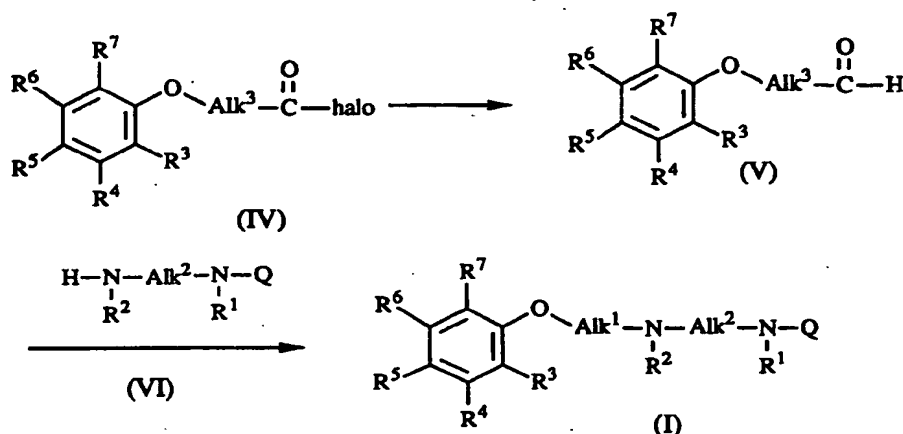


Said reaction can be performed by stirring the diamine of formula (II) with the reagent of formula (III) in an appropriate solvent such as, for example, an alcohol, e.g. ethanol and the like; a halogenated hydrocarbon, e.g. trichloromethane and the like or an ether,

-9-

e.g. tetrahydrofuran and the like; an aromatic hydrocarbon, e.g. methylbenzene and the like; or mixtures thereof. Optionally a base such as, for example, an alkalimetal carbonate, e.g. sodium or potassium carbonate; an alkalimetal hydrogen carbonate, e.g. sodium or potassium hydrogen carbonate; an appropriate organic base, e.g. *N,N*-diethylethanamine, pyridine and the like bases, can be added to pick up the acid that may be formed during the course of the reaction. Elevated temperatures may enhance the rate of the reaction. Preferably the reaction is performed at the reflux temperature of the reaction mixture.

- 10 The compounds of formula (I) can also generally be prepared by reductive *N*-alkylation of an aminoderivative of formula (VI) with an appropriate aldehyde of formula (V), wherein  $\text{Alk}^3$  is  $\text{C}_{1-4}$ alkanediyl.



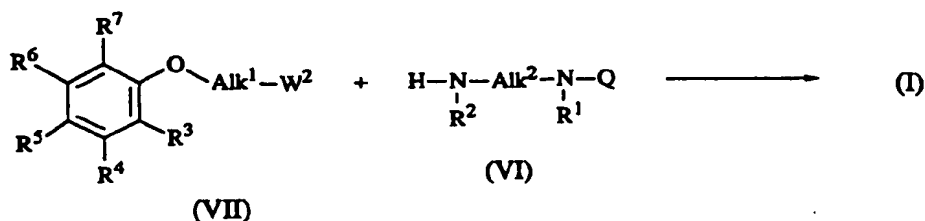
- 15 Said reaction is performed by stirring the reactants of formula (V) and (VI) in an appropriate solvent such as, for example, an alcohol, e.g. ethanol and the like; an ether, e.g. tetrahydrofuran and the like; an aromatic solvent, e.g. methylbenzene and the like, or mixtures thereof. Optionally a water separator can be used to remove the water that is
- 20 formed during the course of the reaction. The resulting imine can then be reduced by reactive hydride reagents such as, for example, sodium borohydride, or by catalytic hydrogenation on an appropriate catalyst such as, for example, palladium on charcoal, platinum on charcoal, Raney nickel and the like in a suitable solvent such as, for example an alcohol, e.g. methanol, ethanol and the like; an ether, e.g. tetrahydrofuran and the
- 25 like; a carboxylic ester, e.g. ethyl acetate, butyl acetate and the like; or a carboxylic acid, e.g. acetic acid, propanoic acid and the like. Optionally the reaction may be performed at elevated temperatures and/or pressures.

The intermediate aldehyde of formula (V) can be prepared by reducing an acyl

-10-

- derivative of formula (IV) wherein  $\text{Alk}^3$  is defined as above. In turn said acyl halide can be prepared by reacting the corresponding, with a halogenating reagent such as thionylchloride, phosphorus trichloride, phosphorus tribromide, oxalylchloride and the like. The latter reaction may be performed in an excess of the halogenating reagent or in appropriate solvents such as, for example, halogenated hydrocarbons, e.g. dichloromethane, trichloromethane and the like; aromatic hydrocarbons, e.g. methylbenzene and the like; ethers, e.g. tetrahydrofuran, 1,4-dioxane and the like, or dipolar aprotic solvents, e.g. N,N-dimethylformamide, N,N-dimethylacetamide and the like. Stirring and elevated temperatures may be appropriate to enhance the rate of the reaction.
- Said reduction of the acylhalide of formula (IV) can for instance be performed by catalytic hydrogenation with a catalyst such as palladium on charcoal, palladium on bariumsulfate, platinum on charcoal and the like in appropriate solvents such as, for example, ethers, e.g. tetrahydrofuran and the like; preferably in admixture with a dipolar aprotic solvent such as, for example, N,N-dimethylformamide, N,N-dimethylacetamide and the like. Optionally a catalyst poison can be added such as thiophene, quinoline-sulfur and the like. The reaction sequence starting from the intermediate of formula (IV) and yielding compounds of formula (I) may be performed as a one-pot procedure.

- The compounds of formula (I) can also be prepared by N-alkylating an amine of formula (VI) with an intermediate of formula (VII), wherein  $\text{W}^2$  is a reactive leaving group such as, for example, halo, e.g. chloro, bromo or iodo; sulfonyloxy, e.g. methanesulfonyloxy, methylbenzenesulfonyloxy and the like, in appropriate solvents such as ketones, e.g. 2-butanone and the like; ethers, e.g. tetrahydrofuran and the like; aromatic hydrocarbons, e.g. methylbenzene and the like; dipolar aprotic solvents, e.g. N,N-dimethylformamide, N,N-dimethylacetamide, dimethylsulfoxide and the like.

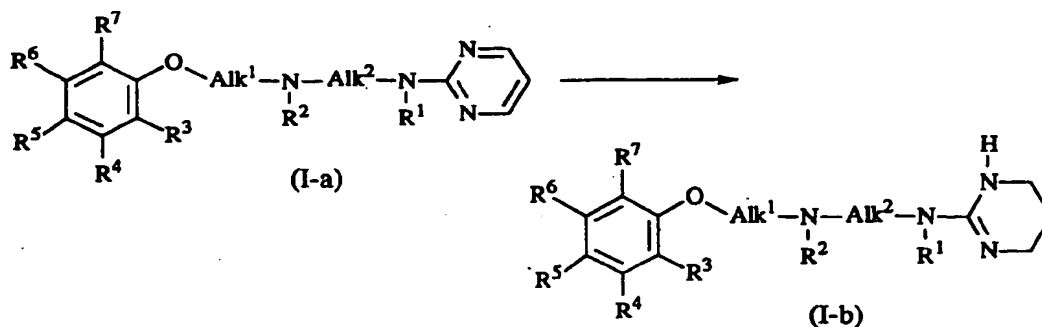


- Stirring and heating may enhance the reaction rate. Optionally a suitable base may be added to pick up the acid that is formed during the course of the reaction such as, for example an alkali metal carbonate, e.g. sodium or potassium carbonate; an alkali metal hydrogen carbonate, e.g. sodium or potassium hydrogen carbonate and the like; an appropriate organic base, e.g. N,N-diethylethanamine, pyridine and the like.

The compounds of formula (I), can also be converted into each other by functional group transformations.

For instance the compounds of formula (I), wherein Q represents a pyrimidinyl moiety can be converted into their tetrahydroanalogs following art-known catalytic

5 hydrogenation procedures.



Furthermore, compounds of formula (I) bearing a C<sub>2</sub>-alkynyl group or C<sub>2</sub>-alkenyl group  
 10 can be converted into the corresponding compounds bearing C<sub>1</sub>-alkyl group following art-known hydrogenation techniques.

Compounds of formula (I) bearing a cyano group can be converted into the corresponding compounds bearing an aminomethyl substituent following art-known hydrogenation techniques.

15 Compounds bearing an alkyloxy substituent can be converted into compounds bearing a hydroxy group by treating the alkyloxy compound with an appropriate acidic reagent such as for example, hydrohalic acid, e.g. hydrobromic acid or borontribromide and the like.

Compounds bearing an arylmethoxy substituent may be converted into compounds  
 20 bearing a hydroxy substituent following art-known debenzylation reactions such as, for example, hydrogenolysis.

Compounds bearing an amino substituent can be N-acylated or N-alkylated following art-known N-acylation or N-alkylation procedures.

Compounds bearing a thio-substituent may be oxidised to the corresponding sulfinyl  
 25 derivatives.

Some of the intermediates mentioned hereinabove are art-known, others are novel and can be prepared following art-known methodologies.

30 Pure stereochemically isomeric forms of the compounds of this invention may be obtained by the application of art-known procedures. Diastereoisomers may be separated

by physical separation methods such as selective crystallization and chromatographic techniques, e.g. liquid chromatography. Enantiomers may be separated from each other by the selective crystallization of their diastereomeric salts with optically active acids. Said pure stereochemically isomeric forms may also be derived from the corresponding  
5 pure stereochemically isomeric forms of the appropriate starting materials, provided that the reaction occurs stereospecifically. Preferably if a specific stereoisomer is desired, said compound will be synthesized by stereospecific methods of preparation. These methods will advantageously employ enantiomerically pure starting materials. Stereochemically isomeric forms of the compounds of formula (I) are obviously intended  
10 to be included within the scope of the invention.

The compounds of formula (I), the pharmaceutically acceptable acid-addition salts and stereochemically isomeric forms thereof have interesting pharmacological properties : they show 5HT<sub>1</sub>-like agonistic activity. The compounds of the present invention have  
15 remarkable vasoconstrictor activity. They are useful to treat conditions which are related to vasodilatation. For instance, they are useful in the treatment of conditions characterized by or associated with cephalic pain, e.g. cluster headache and headache associated with vascular disorders, especially migraine. These compounds are also useful in the treatment of venous insufficiency and in the treatment of conditions  
20 associated with hypotension.

The vasoconstrictor activity of the compounds of formula (I) can be determined using the test described in the pharmacological example, wherein the serotonin-like response of the compounds of the present invention was tested on the basilar arteries of pigs.

25 In view of their useful pharmacological properties, the subject compounds may be formulated into various pharmaceutical forms for administration purposes. To prepare the pharmaceutical compositions of this invention, an effective amount of a particular compound, in base or acid addition salt form, as the active ingredient is combined in intimate admixture with a pharmaceutically acceptable carrier, which carrier  
30 may take a wide variety of forms depending on the form of preparation desired for administration. These pharmaceutical compositions are desirably in unitary dosage form suitable, preferably, for administration orally, rectally, percutaneously, or by parenteral injection. For example, in preparing the compositions in oral dosage form, any of the usual pharmaceutical media may be employed such as, for example, water, glycols, oils,  
35 alcohols and the like in the case of oral liquid preparations such as suspensions, syrups, elixirs and solutions; or solid carriers such as starches, sugars, kaolin, lubricants, binders, disintegrating agents and the like in the case of powders, pills, capsules and tablets. Because of their ease in administration, tablets and capsules represent the most

advantageous oral dosage unit form, in which case solid pharmaceutical carriers are obviously employed. For parenteral compositions, the carrier will usually comprise sterile water, at least in large part, though other ingredients, to aid solubility for example, may be included. Injectable solutions, for example, may be prepared in which the carrier comprises saline solution, glucose solution or a mixture of saline and glucose solution. Injectable suspensions may also be prepared in which case appropriate liquid carriers, suspending agents and the like may be employed. In the compositions suitable for percutaneous administration, the carrier optionally comprises a penetration enhancing agent and/or a suitable wetting agent, optionally combined with suitable additives of any nature in minor proportions, which additives do not cause a significant deleterious effect to the skin. Said additives may facilitate the administration to the skin and/or may be helpful for preparing the desired compositions. These compositions may be administered in various ways, e.g., as a transdermal patch, as a spot-on, as an ointment. It is especially advantageous to formulate the aforementioned pharmaceutical compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used in the specification and claims herein refers to physically discrete units suitable as unitary dosages, each unit containing a predetermined quantity of active ingredient calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. Examples of such dosage unit forms are tablets (including scored or coated tablets), capsules, pills, powder packets, wafers, injectable solutions or suspensions, teaspoonfuls, tablespoonfuls and the like, and segregated multiples thereof.

The compounds of the present invention therefore may be used as medicines in conditions related to vasodilatation, more in particular hypotension, venous insufficiency and especially cephalic pain among which especially migraine. The compounds of the present invention also provide a method of treating warm-blooded animals suffering from conditions related to vasodilatation such as, hypotension, venous insufficiency and especially cephalic pain among which migraine by administering an effective amount of a compound of formula (I), a pharmaceutically acceptable acid addition salt or a stereoisomeric form thereof. Those skilled in the art could easily determine the effective amount from the test results presented hereinafter. In general it is contemplated that an effective amount would be from 1  $\mu\text{g/kg}$  to 1  $\text{mg/kg}$  body weight, and in particular from 2  $\mu\text{g/kg}$  to 200  $\mu\text{g/kg}$  body weight. It may be appropriate to administer the required dose as two, three, four or more sub-doses at appropriate intervals throughout the day. Said sub-doses may be formulated as unit dosage forms, for example, containing 0.005 to 20 mg, and in particular 0.1 mg to 10 mg of active ingredient per unit dosage form.

The following examples are intended to illustrate and not to limit the scope of the

present invention in all its aspects.

### Experimental part

#### A. Preparation of the intermediates

##### 5 Example 1

- a) 2-bromo-1,1-diethoxyethane (0.097mol) was added to a mixture of 2,3-dimethoxyphenol (0.097mol) and potassium carbonate (0.097mol) in *N,N*-dimethylacetamide (200ml). The reaction mixture was stirred for 24 hours at 140°C. The solvent was evaporated. The residue was partitioned between 1,1'-oxybisethane and a solution of
- 10 NaOH in water. The organic layer was separated, washed with a saturated NaCl solution, dried (MgSO<sub>4</sub>), filtered and the solvent was evaporated, yielding 23g (87.7%) of 1-(2,2-diethoxyethoxy)-2,3-dimethoxybenzene (interm. 1).
- b) Hydrochloric acid (2N) (125ml) was added to a solution of intermediate (1) (0.078mol) in 2-propanone (200ml). The reaction mixture was stirred for 15 minutes at
- 15 60°C. The organic solvent was evaporated. Water (300ml) was added. This mixture was extracted with 1,1'-oxybisethane (3x200ml). The separated organic layer was washed with a saturated NaCl solution, dried (MgSO<sub>4</sub>), filtered and the solvent was evaporated, yielding 11.6g (76%) of 2-(2,3-dimethoxyphenoxy)acetaldehyde (interm.2). In a similar manner were also prepared :
- 20 2-[2-(phenylmethoxy)phenoxy]acetaldehyde (interm. 3);  
[2-(methylthio)phenoxy]acetaldehyde (interm. 4); and  
[2-(methylsulfinyl)phenoxy]acetaldehyde (interm. 5).

##### Example 2

- 25 A mixture of *N*-[2-(2-methoxyphenoxy)ethyl]aminepropanenitrile (0.035 mol) in methanol (500ml) was hydrogenated with Raney nickel (2g) as a catalyst. After uptake of hydrogen (2 eq.), the catalyst was filtered off and the filtrate was evaporated, yielding 7.8g (99.4%) of product. A sample (1.0 g) was dissolved in 2-propanol and converted into the hydrochloric acid salt (1:2). The salt was filtered off and dried, yielding 0.81 g
- 30 (60.8%) of *N*-[2-(2-methoxyphenoxy)ethyl]-1,3-propanediamine dihydrochloride; mp. 149.4°C (interm. 6).

##### Example 3

- 35 a) A mixture of 8-methoxy-1,2-benzoxathiin, 2,2-dioxide (0.020 mol) in a hydrobromic acid solution 48% in water (450 ml) was stirred and refluxed for 2 hours. The reaction mixture was cooled. The resulting precipitate was filtered off and the filtrate was extracted with diethyl ether. The separated organic layer was dried (MgSO<sub>4</sub>), filtered and the solvent was evaporated, yielding: 37.4 g 1,2-benzoxathiin-8-ol, 2,2-dioxide (94.4%)



-15-

(interm.7).

- b) A mixture of intermediate 7 (0.13 mol), 2-bromoethanol (0.39 mol) and potassium carbonate (0.015 mol) in 2-propanone (50 ml) was stirred and refluxed overnight. The mixture was cooled and the resulting precipitate was filtered off. The filtrate was  
5 evaporated and the residue was crystallized from  $\text{CH}_2\text{Cl}_2$ . The precipitate was filtered off and the filtrate was evaporated. The residue was purified by column chromatography over silica gel (eluent:  $\text{CH}_2\text{Cl}_2/\text{CH}_3\text{OH}$  97.5/2.5). The pure fractions were collected and the solvent was evaporated, yielding: 3.1 g (9.8%) of 2-(1,2-benzoxathiin-8-yloxy)ethanol 2,2-dioxide (interm.8)
- 10 c) *N,N*,diethylethanamine (10 ml) was added dropwise to a mixture of intermediate 8 (0.089 mol) and methanesulfonyl chloride (0.13 mol) in 2-propanone (250 ml), stirred and cooled on an ice bath. The reaction mixture was stirred for 1 hour at room temperature. The mixture was filtered and the filtrate was evaporated. The residue was dissolved in  $\text{CH}_2\text{Cl}_2$ . The organic solution was washed with an aqueous hydrochloric  
15 acid solution, dried ( $\text{MgSO}_4$ ), filtered and the solvent was evaporated. The residue was crystallized from  $\text{CH}_2\text{Cl}_2$ . The precipitate was filtered off and dried (vacuum; 70 °C), yielding 15.6 g (54.7%) 2-(1,2-benzoxathiin-8-yloxy)ethanol 2,2-dioxide methanesulfonate (ester); mp. 117 °C (interm. 9).
- d) A mixture of intermediate 9 (0.019 mol) in methanol (250 ml) was hydrogenated with  
20 palladium-on-charcoal catalyst (2 g) as a catalyst. After uptake of hydrogen ( $\text{H}_2$ ) (1 equiv), the catalyst was filtered off and the filtrate was evaporated. The residue was crystallized from  $\text{CH}_3\text{CN}$ . The precipitate was filtered off and dried (vacuum; 60 °C), yielding: 3.1 g (50.6%) 2-[(3,4-dihydro-1,2-benzoxathiin-8-yl)oxy]ethanol methanesulfonate(ester) 2,2-dioxide; mp. 155 °C (interm.10)

25

#### Example 4

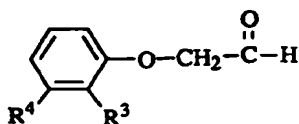
- a) A mixture of 2,3-dihydro-5-hydroxy-1,4-benzodioxin (0.13mol), 2-bromo-1,1-diethoxyethane (0.13mol) and potassium carbonate (0.13mol) in *N,N*-dimethylacetamide (250ml) was stirred overnight at 140 °C. The solvent was evaporated. The  
30 residue was partitioned between 1,1'-oxybisethane and water. The organic layer was separated, washed with a saturated NaCl solution, dried ( $\text{MgSO}_4$ ), filtered and the solvent was evaporated. The residual oil was crystallized from 2,2'-oxybispropane. The precipitate was filtered off and dried, yielding 21g (60.2%) 5-(2,2-diethoxyethoxy)-2,3-dihydro-1,4-benzodioxin; mp. 73.1 °C (interm.11).
- 35 b) Hydrochloric acid 2N (125ml) was added to a solution of intermediate (R 97.205) (0.078mol) in 2-propanone (200ml). The reaction mixture was stirred for 15 minutes at 60 °C. The organic solvent was evaporated (40 °C). Water (300ml) was added. This

-16-

mixture was extracted with dichloromethane. The separated organic layer was dried (MgSO<sub>4</sub>), filtered and the solvent was evaporated, yielding 13g (86.6%) of [(2,3-dihydro-1,4-benzodioxin-5-yl)oxy]acetaldehyde (interm.12).

# 5 Table 1.

In this manner were prepared :



10

Int. no.	R <sup>3</sup> , R <sup>4</sup>
12	-O-(CH <sub>2</sub> ) <sub>2</sub> -O-
13	-(CH <sub>2</sub> ) <sub>4</sub> -
14	-O-CH=CH-
15	-O-(CH <sub>2</sub> ) <sub>3</sub> -
16	-(CH <sub>2</sub> ) <sub>3</sub> -O-
17	-C(CH <sub>3</sub> )=C(CH <sub>3</sub> )-O-

## B. Preparation of the final compounds

### Example 5

A mixture of intermediate 6 (0.03mol), 2-chloropyrimidine (0.03mol) and sodium carbonate (0.03mol) in ethanol (150ml) was stirred and refluxed overnight. The reaction mixture was filtered over dicalite. The filtrate was evaporated. The residue was dissolved in acetonitrile and this mixture was acidified with HCl/2-propanol. The precipitate was filtered off. The filtrate was evaporated and the residue was stirred in water. This mixture was alkalized with NaOH, then extracted with 1,1'-oxybisethane. The separated organic layer was dried (MgSO<sub>4</sub>), filtered and the solvent was evaporated. The residue was dissolved in warm methanol (500ml) and converted into the ethanedioic acid salt (1:2) with a solution of ethanedioic acid (8g) in methanol. The salt was filtered off and dried, yielding 6.8g (56.3%) of N-[2-(2-methoxyphenoxy)ethyl]-N'-2-pyrimidinyl-1,3-propanediamine ethanedioate (1:2); mp. 178.4°C (comp. 1).

25

### Example 6

N-2-pyrimidinyl-1,3-propanediamine (0.042mol) was added to a solution of intermediate 2 (0.056mol) in ethanol (200ml) and this mixture was stirred for 30 min. at room

-17-

temperature. The reaction mixture was cooled to 0°C with an ice salt bath. Sodium borohydride (0.059 mol) was added and the reaction mixture was stirred for 30 minutes at 0°C, then for 1 hour at room temperature. A small amount of water was added and the solvent was evaporated at 40°C. The residue was partitioned between dichloromethane and water. The organic layer was separated, dried (MgSO<sub>4</sub>), filtered and the solvent was evaporated. The residual oil (13 g) was purified by column chromatography over silica gel (eluent: CH<sub>2</sub>Cl<sub>2</sub>/(CH<sub>3</sub>OH/NH<sub>3</sub>) 95/5). The desired fractions were collected and the solvent was evaporated. The residual oil (8 g) was dissolved in 2-propanone and converted into the ethanedioic acid salt (1:1). The precipitate was filtered off and crystallized from methanol. The precipitate was filtered off and dried, yielding 6.7 g (37.8%) of N-[2-(2,3-dimethoxyphenoxy)ethyl]-N'-2-pyrimidinyl-1,3-propanediamine ethanedioate(1:1); mp. 200.0°C (comp. 2).

#### Example 7

- 15 a) A mixture of intermediate 3 (0.117 mol) and N-2-pyrimidinyl-1,3-propanediamine (0.087 mol) in ethanol (500 ml) was stirred for 45 minutes at 20°C. The reaction mixture was cooled to 0°C (ice salt bath). Sodium borohydride (0.125 mol) was added in one portion and the reaction mixture was stirred for 2 hours. Water was added and the solvent was evaporated. The residue was partitioned between 1,1'-oxybisethane and water. The organic layer was separated, dried, filtered and the solvent was evaporated. The residue was purified by column chromatography over silica gel (eluent: CH<sub>2</sub>Cl<sub>2</sub>/(CH<sub>3</sub>OH/NH<sub>3</sub>) 95/5). The pure fractions were collected and the solvent was evaporated. The residue was crystallized from 2-propanol. The precipitate was filtered off and dried, yielding 7.7 g (23.4%) of N-[2-[2-(phenylmethoxy)phenoxy]ethyl]-N'-2-pyrimidinyl-1,3-propanediamine; mp. 90.1°C (comp. 3).
- 25 b) A mixture of compound (97232) (0.02 mol) in methanol (250 ml) was hydrogenated with palladium on activated carbon 10% (2 g) as a catalyst. After uptake of hydrogen (1 eq.), the catalyst was filtered off. The filtrate was evaporated. The residue was dissolved in ethanol and converted into the hydrochloric acid salt (1:2) with HCl/2-propanol. The salt was filtered off and dried, yielding 4.8 g (66.4%) of 2-[2-[[3-(2-pyrimidinylamino)propyl]amino]ethoxy]phenol dihydrochloride; mp. 166.4°C (comp. 4).

#### Example 8

- 35 A mixture of 1-bromo-3-(2-methoxyphenoxy)propane (0.020 mol), N-2-pyrimidinyl-1,3-propanediamine (0.020 mol) and potassium carbonate (0.03 mol) in N,N-dimethylacetamide (50 ml) was stirred for 48 hours at 70 °C. The solvent was evaporated. The residue was partitioned between dichloromethane and water. The organic layer was

separated, dried (MgSO<sub>4</sub>), filtered and the solvent was evaporated. The residual oil was purified by column chromatography over silica gel (eluent: CH<sub>2</sub>Cl<sub>2</sub>/(CH<sub>3</sub>OH/NH<sub>3</sub>) 95/5). The desired fractions were collected and the solvent was evaporated. The residue (5 g) was dissolved in 2-propanone and converted into the ethanedioic acid salt (1:1).

- 5 The precipitate was filtered off and crystallized from methanol. The precipitate was filtered off and dried, yielding 3.41g of product. This fraction was recrystallized from methanol. The precipitate was filtered off and dried, yielding 3.3 g (40.6%) N-[3-(2-methoxyphenoxy)propyl]-N'-2-pyrimidinyl-1,3-propanediamine ethanedioate(1:1); mp. 186.3°C (comp. 5).

10

#### Example 9

A mixture of compound 2 (0.0135mol) and ethanedioic acid dihydrate (0.0135mol) in 2-methoxyethanol (300ml) was hydrogenated at 80°C with palladium on activated carbon 10% (2g) as a catalyst in the presence of a 4% solution of thiophene (2ml). After uptake of hydrogen (2 eq.), the catalyst was filtered off and the filtrate was evaporated. The residue was crystallized from methanol. The precipitate was filtered off and dried, yielding 2.56g (36.7%) of N-[2-(2,3-dimethoxyphenoxy)ethyl]-N'-(1,4,5,6-tetrahydro-2-pyrimidinyl)-1,3-propanediamine ethanedioate (1:2); mp. 181.1°C (comp. 6).

#### 20 Example 10

N-2-pyrimidinyl-1,3-propanediamine (0.05mol) was added to a solution of intermediate 12 (0.067mol) in ethanol (200ml) and this mixture was stirred for 30 minutes at room temperature. The reaction mixture was cooled to 0°C with an ice salt bath. Sodium borohydride (0.070mol) was added and the reaction mixture was stirred for 30 minutes at 0°C, then for 30 minutes at room temperature. A small amount of water was added and the solvent was evaporated. The residue was partitioned between dichloromethane and water. The organic layer was separated, dried (MgSO<sub>4</sub>), filtered and the solvent was evaporated. The residual oil was purified by column chromatography over silica gel (eluent: CH<sub>2</sub>Cl<sub>2</sub>/(CH<sub>3</sub>OH/NH<sub>3</sub>) 95/5). The desired fractions were collected and the solvent was evaporated. The residual oil was dissolved in 2-propanone and converted into the ethanedioic acid salt (1:2). The precipitate was filtered off and crystallized from methanol. The precipitate was filtered off and dried, yielding 6g (23.5%) of N-[2-[(2,3-dihydro-1,4-benzodioxin-5-yl)oxy]ethyl]-N'-2-pyrimidinyl-1,3-propanediamine ethanedioate (1:1); mp. 213.2°C (comp. 7).

35

#### Example 11

A mixture of 5-(3-chloropropoxy)-2,3-dihydro-1,4-benzodioxin (0.017 mol), N-2-pyrimidinyl-1,3-propanediamine (0.026 mol) and calcium oxide (5 g) in tetrahydro-

5 furan (150 ml) was stirred overnight at 160 °C (pressure vessel). The mixture was cooled, filtered and the filtrate was evaporated. The residue was purified by column chromatography over silica gel (eluent: CH<sub>2</sub>Cl<sub>2</sub>/(CH<sub>3</sub>OH/NH<sub>3</sub>) 95/5). The pure fractions were collected and the solvent was evaporated, yielding 2.66 g N-[3-[(2,3-dihydro-1,4-benzodioxin-5-yl)oxy]propyl]-N'-2-pyrimidinyl-1,3-propanediamine ethanedioate(1:1) (36.0%); mp. 200.2°C (comp. 8)

#### Example 12

10 Compound 7 (0.0078mol) and ethanedioic acid dihydrate (0.0078mol) were dissolved in a warm mixture of 2-methoxyethanol (200ml) and water (100ml). This solution was hydrogenated at 80°C with palladium on activated carbon (10%) (2g) as a catalyst in the presence of a 4% thiophene solution (1ml). After uptake of hydrogen (2 eq.), the catalyst was filtered off and the filtrate was evaporated. The residue was crystallized from methanol. The precipitate was filtered off and dried. This fraction was  
15 recrystallized from water. The precipitate was filtered off and dried, yielding 0.8g of product. This fraction was recrystallized from methanol/water (5/1). The precipitate was filtered off and dried, yielding 0.5g (13.7%) of N-[2-[(2,3-dihydro-1,4-benzodioxin-5-yl)oxy]ethyl]-N'-(1,4,5,6-tetrahydro-2-pyrimidinyl)-1,3-propanediamine ethanedioate(2:3); mp. 231.1°C (comp.9 ).

#### Example 13

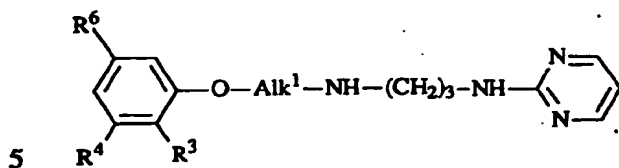
[phenoxy-1-(1methylethyl)methylene]cyanamide (0.019 mol) was added to a solution of N-[(2,3-dihydro-1,4-benzodioxin-5-yl)oxy]ethyl]-1,4-propanediamine (0.019 mol) in methanol (100 ml). The reaction mixture was stirred for 4 days at room temperature.  
25 The solvent was evaporated. The resultant oil was purified by column chromatography over silica gel (eluent: CH<sub>2</sub>Cl<sub>2</sub>/CH<sub>3</sub>OH 90/10). The pure fractions were collected and the solvent was evaporated. The resultant oil (2.9 g) was crystallized from CH<sub>3</sub>CN. The precipitate was filtered off and dried, yielding: 2.4 g (34.9%) N"-cyano-N-[3-[[2-[(2,3-dihydro-1,4-benzodioxin-5-yl)oxy]ethyl]amino]propyl]-N'-(1-methylethyl)guanidine;  
30 mp. 120.6°C (comp.10).

#### Example 14

A mixture of compound 10 (0.003 mol) in hydrochloric acid in 2-propanol (10 ml) and methanol (50 ml) was stirred and refluxed for 30 minutes. The solvent was evaporated.  
35 The residue was crystallized from CH<sub>3</sub>CN. The precipitate was filtered off and dried, yielding: 0.53 g (39.1%) of N-[[[3-[[2-[(2,3-dihydro-1,4-benzodioxin-5-yl)oxy]ethyl]amino]propyl]amino][(1-methylethyl)amino]methylene]urea dihydrochloride; mp. 155.0°C (comp.11).

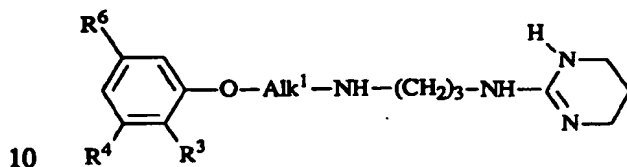
In this manners were prepared :

Table 2



Co.No.	Ex. No.	Alk <sup>1</sup>	R <sup>3</sup>	R <sup>4</sup> , R <sup>6</sup>	physical data
1	5	(CH <sub>2</sub> ) <sub>2</sub>	O-CH <sub>3</sub>	H	mp. 178.4°C / . 2 (COOH) <sub>2</sub>
2	6	(CH <sub>2</sub> ) <sub>2</sub>	O-CH <sub>3</sub>	3-O-CH <sub>3</sub>	mp. 200.0°C / . (COOH) <sub>2</sub>
3	7a	(CH <sub>2</sub> ) <sub>2</sub>	O-CH <sub>2</sub> -C <sub>6</sub> H <sub>5</sub>	H	mp. 90.1°C
4	7b	(CH <sub>2</sub> ) <sub>2</sub>	OH	H	mp. 166.4°C / . 2 HCl
5	8	(CH <sub>2</sub> ) <sub>3</sub>	O-CH <sub>3</sub>	H	mp. 186.3°C / . (COOH) <sub>2</sub>
12	6	CH(CH <sub>3</sub> )CH <sub>2</sub>	O-CH <sub>3</sub>	H	mp. 167.9°C / . 2 HCl
13	6	(CH <sub>2</sub> ) <sub>2</sub>	CH <sub>3</sub>	H	mp. 204.8°C / . (COOH) <sub>2</sub>
14	6	(CH <sub>2</sub> ) <sub>2</sub>	O-CH <sub>2</sub> -CH <sub>3</sub>	H	mp. 160.3°C / . 2 (COOH) <sub>2</sub>
15	6	(CH <sub>2</sub> ) <sub>2</sub>	O-CH <sub>3</sub>	5-CH <sub>3</sub>	mp. 197.2°C / . (COOH) <sub>2</sub>
16	8	(CH <sub>2</sub> ) <sub>2</sub>	CO-CH <sub>3</sub>	H	mp. 179.0°C / (COOH) <sub>2</sub>
17	6	(CH <sub>2</sub> ) <sub>2</sub>	S-CH <sub>3</sub>	H	mp. 217.6°C / (COOH) <sub>2</sub>
18	8	(CH <sub>2</sub> ) <sub>2</sub>	CN	H	mp. 185.1°C / (COOH) <sub>2</sub>
19	6	(CH <sub>2</sub> ) <sub>2</sub>	SO-CH <sub>3</sub>	H	mp. 177.7°C / 2 (COOH) <sub>2</sub>
20	8	(CH <sub>2</sub> ) <sub>2</sub>	Br	H	mp. 198.2° / (COOH) <sub>2</sub>

Table 3

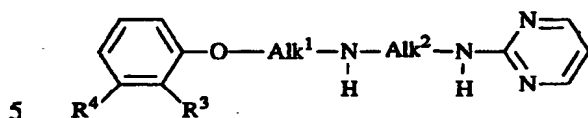


Co. No.	Ex. No.	Alk <sup>1</sup>	R <sup>3</sup>	R <sup>4</sup> , R <sup>6</sup>	physical data
6	9	(CH <sub>2</sub> ) <sub>2</sub>	O-CH <sub>3</sub>	3-O-CH <sub>3</sub>	mp. 181.1°C / . 2 (COOH) <sub>2</sub>
21	9	(CH <sub>2</sub> ) <sub>2</sub>	CH <sub>3</sub>	H	mp. 170.3°C / . 2 HCl
22	9	(CH <sub>2</sub> ) <sub>2</sub>	O-CH <sub>3</sub>	H	mp. 159.1°C / . 2 (COOH) <sub>2</sub>
23	9	(CH <sub>2</sub> ) <sub>2</sub>	O-CH <sub>2</sub> -CH <sub>3</sub>	H	mp. 168.1°C / . 2 (COOH) <sub>2</sub>

-21-

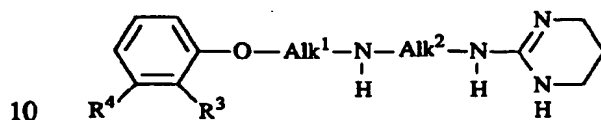
Co. No.	Ex. No.	Alk <sup>1</sup>	R <sup>3</sup>	R <sup>4</sup> , R <sup>6</sup>	physical data
24	9	(CH <sub>2</sub> ) <sub>2</sub>	O-CH <sub>3</sub>	5-CH <sub>3</sub>	mp. 182.8°C / . 2 (COOH) <sub>2</sub>
25	9	(CH <sub>2</sub> ) <sub>2</sub>	OH	H	mp. 185.4°C / . 2 HCl
26	9	(CH <sub>2</sub> ) <sub>3</sub>	O-CH <sub>3</sub>	H	mp. 155.1°C / . 2 (COOH) <sub>2</sub>
27	9	(CH <sub>2</sub> ) <sub>2</sub>	CO-CH <sub>3</sub>	H	mp. 150.6°C / 2 (COOH) <sub>2</sub> ½ H <sub>2</sub> O
28	9	(CH <sub>2</sub> ) <sub>2</sub>	CN	H	mp. 188.6°C / 3/2 (COOH) <sub>2</sub>

Table 4



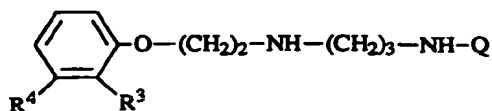
Co. No.	Ex. No	R <sup>3</sup> , R <sup>4</sup>	Alk <sup>1</sup>	Alk <sup>2</sup>	Physical data
7	10	-O-(CH <sub>2</sub> ) <sub>2</sub> -O-	(CH <sub>2</sub> ) <sub>2</sub>	(CH <sub>2</sub> ) <sub>3</sub>	mp. 213.2°C / . (COOH) <sub>2</sub>
29	10	-O-(CH <sub>2</sub> ) <sub>2</sub> -O-	(CH <sub>2</sub> ) <sub>2</sub>	(CH <sub>2</sub> ) <sub>4</sub>	mp. 210.1°C / . (COOH) <sub>2</sub>
30	10	-O-(CH <sub>2</sub> ) <sub>2</sub> -O-	(CH <sub>2</sub> ) <sub>2</sub>	(CH <sub>2</sub> ) <sub>2</sub>	mp. 204.1°C / . (COOH) <sub>2</sub>
31	10	-CH=CH-CH=CH-	(CH <sub>2</sub> ) <sub>2</sub>	(CH <sub>2</sub> ) <sub>3</sub>	mp. 227.6°C / . (COOH) <sub>2</sub>
32	10	-(CH <sub>2</sub> ) <sub>4</sub> -	(CH <sub>2</sub> ) <sub>2</sub>	(CH <sub>2</sub> ) <sub>3</sub>	mp. 229.9°C / . (COOH) <sub>2</sub>
33	10	-O-CH=CH-	(CH <sub>2</sub> ) <sub>2</sub>	(CH <sub>2</sub> ) <sub>3</sub>	mp. 223.3°C / . (COOH) <sub>2</sub>
34	10	-O-(CH <sub>2</sub> ) <sub>3</sub> -	(CH <sub>2</sub> ) <sub>2</sub>	(CH <sub>2</sub> ) <sub>3</sub>	mp. 206.6°C / . (COOH) <sub>2</sub>
8	11	-O-(CH <sub>2</sub> ) <sub>2</sub> -	(CH <sub>2</sub> ) <sub>3</sub>	(CH <sub>2</sub> ) <sub>3</sub>	mp. 200.0°C / . (COOH) <sub>2</sub>
35	10	-S-CH=CH-	(CH <sub>2</sub> ) <sub>2</sub>	(CH <sub>2</sub> ) <sub>3</sub>	mp. 227.2°C / . (COOH) <sub>2</sub>
36	10	-(CH <sub>2</sub> ) <sub>3</sub> -O-	(CH <sub>2</sub> ) <sub>2</sub>	(CH <sub>2</sub> ) <sub>3</sub>	mp. 67.8°C
37	10	-(CH <sub>2</sub> ) <sub>3</sub> -O-	(CH <sub>2</sub> ) <sub>2</sub>	(CH <sub>2</sub> ) <sub>3</sub>	mp. 219°C / . (COOH) <sub>2</sub>
38	10	-C(CH <sub>3</sub> )=C(CH <sub>3</sub> )-O-	(CH <sub>2</sub> ) <sub>2</sub>	(CH <sub>2</sub> ) <sub>2</sub>	mp. 87.1°C
39	11	-(CH <sub>2</sub> ) <sub>2</sub> -S(O) <sub>2</sub> -O-	(CH <sub>2</sub> ) <sub>2</sub>	(CH <sub>2</sub> ) <sub>2</sub>	mp. 207.5°C / . (COOH) <sub>2</sub>

Table 5



Co. No.	Ex. No.	R <sup>3</sup> , R <sup>4</sup>	Alk <sup>1</sup>	Alk <sup>2</sup>	Physical data
9	12	-O-(CH <sub>2</sub> ) <sub>2</sub> -O	-(CH <sub>2</sub> ) <sub>2</sub> -	-(CH <sub>2</sub> ) <sub>3</sub> -	mp. 231.1°C/ . 3/2 (COOH) <sub>2</sub>
40	12	-O-(CH <sub>2</sub> ) <sub>2</sub> -O	-(CH <sub>2</sub> ) <sub>2</sub> -	-(CH <sub>2</sub> ) <sub>4</sub> -	mp. 193.7°C/ . 2 (COOH) <sub>2</sub>
41	12	-O-(CH <sub>2</sub> ) <sub>2</sub> -O	-(CH <sub>2</sub> ) <sub>2</sub> -	-(CH <sub>2</sub> ) <sub>2</sub> -	mp. 213.5°C/ . 2 (COOH) <sub>2</sub>
42	12	-CH=CH-CH=CH-	-(CH <sub>2</sub> ) <sub>2</sub> -	-(CH <sub>2</sub> ) <sub>3</sub> -	mp. 221.3°C/ . 2 (COOH) <sub>2</sub>
43	12	-(CH <sub>2</sub> ) <sub>4</sub> -	-(CH <sub>2</sub> ) <sub>2</sub> -	-(CH <sub>2</sub> ) <sub>3</sub> -	mp. 205.6°C/ . 2 (COOH) <sub>2</sub>
44	12	-O-(CH <sub>2</sub> ) <sub>3</sub> -	-(CH <sub>2</sub> ) <sub>2</sub> -	-(CH <sub>2</sub> ) <sub>3</sub> -	mp. 230.8°C/ . 3/2 (COOH) <sub>2</sub>
45	12	-O-CH=CH-	-(CH <sub>2</sub> ) <sub>2</sub> -	-(CH <sub>2</sub> ) <sub>3</sub> -	2 (COOH) <sub>2</sub>
46	12	-O-(CH <sub>2</sub> ) <sub>2</sub> -O-	-(CH <sub>2</sub> ) <sub>3</sub> -	-(CH <sub>2</sub> ) <sub>3</sub> -	mp. 205.5°C/ . 2 (COOH) <sub>2</sub>
47	12	-(CH <sub>2</sub> ) <sub>3</sub> -O-	-(CH <sub>2</sub> ) <sub>2</sub> -	-(CH <sub>2</sub> ) <sub>3</sub> -	mp. 191.1°C/ . 2 (COOH) <sub>2</sub>
48	12	-C(CH <sub>3</sub> )=C(CH <sub>3</sub> )O-	-(CH <sub>2</sub> ) <sub>2</sub> -	-(CH <sub>2</sub> ) <sub>3</sub> -	mp. 194.0°C/ . 2 (COOH) <sub>2</sub>

Table 6



Co. No.	Ex. No.	R <sup>3</sup> , R <sup>4</sup>	Q	physical data
10	13	-O-(CH <sub>2</sub> ) <sub>2</sub> -O-	$\begin{array}{c} \text{N} \text{---} \text{CN} \\   \\ \text{---} \text{NH} \text{---} \text{CH}(\text{CH}_3)_2 \end{array}$	mp. 181.1°C/ . 2 (COOH) <sub>2</sub>
11	14	-O-(CH <sub>2</sub> ) <sub>2</sub> -O-	$\begin{array}{c} \text{O} \\    \\ \text{N} \text{---} \text{C} \text{---} \text{NH}_2 \\   \\ \text{---} \text{NH} \text{---} \text{CH}(\text{CH}_3)_2 \end{array}$	mp. 155.0°C/ . 2 HCl

### C. Pharmacological Example

#### Example 15

10 Segments of basilar arteries taken from pigs (anaesthetised with sodium pentobarbital) were mounted for recording of isometric tension in organ baths. The preparations were bathed in Krebs - Henseleit solution. The solution was kept at 37°C and gassed with a mixture of 95% O<sub>2</sub> - 5% CO<sub>2</sub>. The preparations were stretched until a stable basal tension of 2 grams was obtained.

15 The preparations were made to constrict with serotonin (3x10<sup>-7</sup> M). The response to the addition of serotonin was measured and subsequently the serotonin was washed away. This procedure was repeated until stable responses were obtained.

Subsequently the test compound was administered to the organ bath and the constriction



of the preparation was measured. This constrictive response was expressed as a percentage of the response to serotonin as measured previously.

The lowest active concentration was defined as the concentration at which 50% of the response to serotonin is obtained.

5

In table 7 the lowest active concentration of compounds of formula (I) are presented.

Table 7

Co. No.	lowest active concentration (M)
1	$1 \cdot 10^{-6}$
2	$3 \cdot 10^{-7}$
4	$1 \cdot 10^{-6}$
6	$1 \cdot 10^{-6}$
7	$3 \cdot 10^{-7}$
9	$3 \cdot 10^{-8}$
10	$3 \cdot 10^{-7}$
11	$1 \cdot 10^{-7}$
16	$1 \cdot 10^{-7}$
17	$1 \cdot 10^{-6}$
18	$1 \cdot 10^{-6}$
19	$1 \cdot 10^{-6}$
20	$1 \cdot 10^{-7}$
21	$1 \cdot 10^{-6}$
22	$1 \cdot 10^{-6}$
23	$3 \cdot 10^{-7}$
25	$1 \cdot 10^{-6}$
26	$3 \cdot 10^{-7}$
27	$1 \cdot 10^{-7}$
29	$1 \cdot 10^{-6}$
31	$3 \cdot 10^{-8}$
32	$3 \cdot 10^{-7}$
35	$3 \cdot 10^{-7}$
36	$3 \cdot 10^{-8}$
38	$1 \cdot 10^{-6}$
39	$1 \cdot 10^{-6}$
40	$3 \cdot 10^{-6}$

Co. No.	lowest active concentration (M)
44	$1 \cdot 10^{-6}$
45	$3 \cdot 10^{-7}$
47	$1 \cdot 10^{-7}$
48	$1 \cdot 10^{-6}$

**D. Composition examples**

- “Active ingredient” (A.I.) as used throughout these examples relates to a compound of formula (I), a pharmaceutically acceptable acid addition salt or a stereochemically isomeric form thereof.

**Example 16 : ORAL DROPS**

- 500 Grams of the A.I. was dissolved in 0.5 l of 2-hydroxypropanoic acid and 1.5 l of the polyethylene glycol at 60–80°C. After cooling to 30–40°C there were added 35 l of polyethylene glycol and the mixture was stirred well. Then there was added a solution of 1750 grams of sodium saccharin in 2.5 l of purified water and while stirring there were added 2.5 l of cocoa flavor and polyethylene glycol q.s. to a volume of 50 l, providing an oral drop solution comprising 10 mg/ml of A.I.. The resulting solution was filled into suitable containers.

**Example 17 : ORAL SOLUTION**

- 9 Grams of methyl 4-hydroxybenzoate and 1 gram of propyl 4-hydroxybenzoate were dissolved in 4 l of boiling purified water. In 3 l of this solution were dissolved first 10 grams of 2,3-dihydroxybutanedioic acid and thereafter 20 grams of the A.I. The latter solution was combined with the remaining part of the former solution and 12 l 1,2,3-propanetriol and 3 l of sorbitol 70% solution were added thereto. 40 Grams of sodium saccharin were dissolved in 0.5 l of water and 2 ml of raspberry and 2 ml of gooseberry essence were added. The latter solution was combined with the former, water was added q.s. to a volume of 20 l providing an oral solution comprising 5 mg of the active ingredient per teaspoonful (5 ml). The resulting solution was filled in suitable containers.

**Example 18 : CAPSULES**

- 20 Grams of the A.I., 6 grams sodium lauryl sulfate, 56 grams starch, 56 grams lactose, 0.8 grams colloidal silicon dioxide, and 1.2 grams magnesium stearate were vigorously stirred together. The resulting mixture was subsequently filled into 1000 suitable hardened gelatin capsules, comprising each 20 mg of the active ingredient.

**Example 19 : FILM-COATED TABLETS****Preparation of tablet core**

A mixture of 100 grams of the A.I., 570 grams lactose and 200 grams starch was mixed well and thereafter humidified with a solution of 5 grams sodium dodecyl sulfate and 10 grams polyvinylpyrrolidone in about 200 ml of water. The wet powder mixture was sieved, dried and sieved again. Then there was added 100 grams microcrystalline cellulose and 15 grams hydrogenated vegetable oil. The whole was mixed well and compressed into tablets, giving 10.000 tablets, each containing 10 mg of the active ingredient.

**10 Coating**

To a solution of 10 grams methyl cellulose in 75 ml of denaturated ethanol there was added a solution of 5 grams of ethyl cellulose in 150 ml of dichloromethane. Then there were added 75 ml of dichloromethane and 2.5 ml 1,2,3-propanetriol. 10 Grams of polyethylene glycol was molten and dissolved in 75 ml of dichloromethane. The latter solution was added to the former and then there were added 2.5 grams of magnesium octadecanoate, 5 grams of polyvinylpyrrolidone and 30 ml of concentrated colour suspension and the whole was homogenated. The tablet cores were coated with the thus obtained mixture in a coating apparatus.

**Example 20 : INJECTABLE SOLUTION**

1.8 Grams methyl 4-hydroxybenzoate and 0.2 grams propyl 4-hydroxybenzoate were dissolved in about 0.5 l of boiling water for injection. After cooling to about 50°C there were added while stirring 4 grams lactic acid, 0.05 grams propylene glycol and 4 grams of the A.I.. The solution was cooled to room temperature and supplemented with water for injection q.s. ad 1 l, giving a solution comprising 4 mg/ml of A.I.. The solution was sterilized by filtration (U.S.P. XVII p. 811) and filled in sterile containers.

**Example 21 : SUPPOSITORIES**

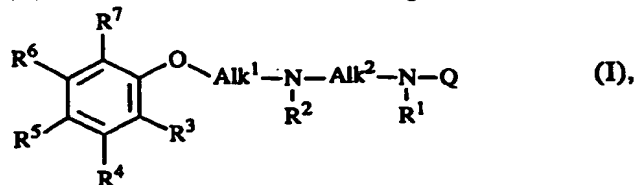
3 Grams A.I. was dissolved in a solution of 3 grams 2,3-dihydroxybutanedioic acid in 25 ml polyethylene glycol 400. 12 Grams surfactant (SPAN®) and triglycerides (Witepsol 555 ®) q.s. ad 300 grams were molten together. The latter mixture was mixed well with the former solution. The thus obtained mixture was poured into moulds at a temperature of 37-38°C to form 100 suppositories each containing 30 mg/ml of the A.I.

**Example 22 : INJECTABLE SOLUTION**

60 Grams of A.I. and 12 grams of benzylalcohol were mixed well and sesame oil was added q.s. ad 1 l, giving a solution comprising 60 mg/ml of A.I. The solution was sterilized and filled in sterile containers.

# Claims

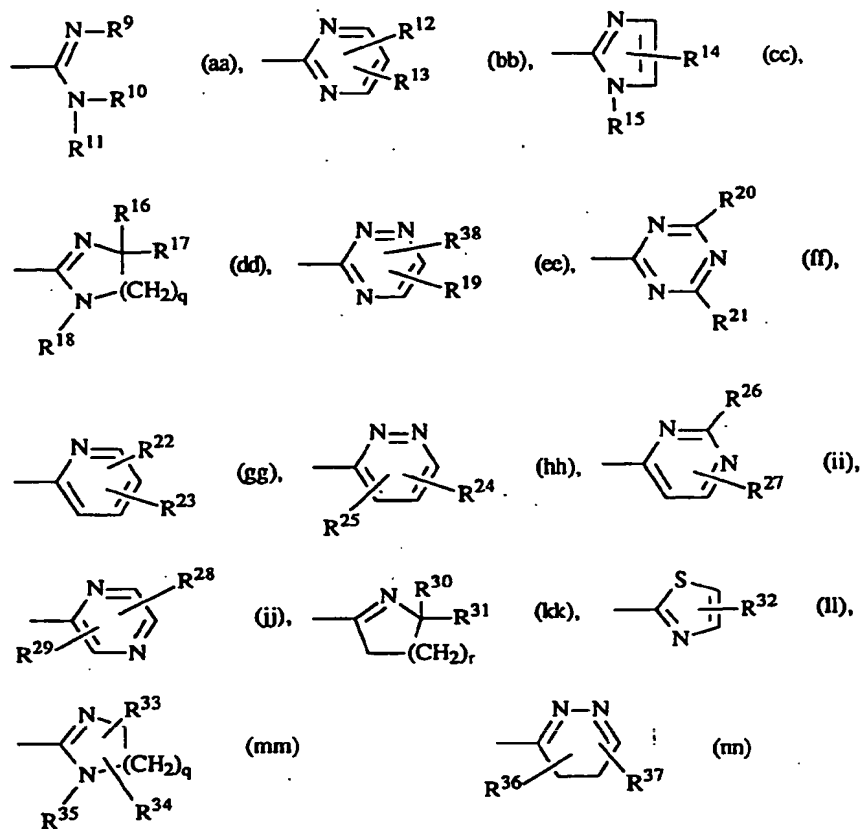
1. Use of a compound for the manufacture of a medicament for the treatment of conditions which are related to vasodilatation, having the formula



the pharmaceutically acceptable acid addition salts thereof, and the stereochemically isomeric forms thereof, wherein

- 10  $\text{R}^1$  and  $\text{R}^2$  each independently are hydrogen or  $\text{C}_{1-6}$ alkyl;  
 $\text{R}^3$  is  $\text{C}_{1-6}$ alkyl, hydroxy, cyano, halo,  $\text{C}_{1-6}$ alkyloxy, aryloxy, arylmethoxy,  $\text{C}_{2-6}$ alkenyl,  $\text{C}_{2-6}$ alkynyl,  $\text{C}_{1-6}$ alkyl-S-,  $\text{C}_{1-6}$ alkyl(S=O)-,  $\text{C}_{1-6}$ alkylcarbonyl;  
 $\text{R}^4$  is hydrogen, halo, hydroxy,  $\text{C}_{1-6}$ alkyl, or  $\text{C}_{1-6}$ alkyloxy; or  $\text{R}^3$  and  $\text{R}^4$  taken together form a bivalent radical of formula
- |  |  |
|--|--|
| $\text{---CH=CH---CH=CH---}$ (a),              | $\text{---X---CH=CH---}$ (f),                      |
| $\text{---(CH}_2\text{)}_n\text{---}$ (b),     | $\text{---O---(CH}_2\text{)}_t\text{---Y---}$ (g), |
| $\text{---(CH}_2\text{)}_m\text{---X---}$ (c), | $\text{---Y---(CH}_2\text{)}_t\text{---O---}$ (h), |
| $\text{---X---(CH}_2\text{)}_m\text{---}$ (d), | $\text{---(CH}_2\text{)}_t\text{---Z---}$ (i),     |
| $\text{---CH=CH---X---}$ (e),                  | $\text{---Z---(CH}_2\text{)}_t\text{---}$ (j),     |
- 15 in these bivalent radicals one or two hydrogen atoms may be substituted with  $\text{C}_{1-6}$ alkyl,  $\text{C}_{1-6}$ alkylcarbonyl or  $\text{C}_{1-6}$ alkyl-S(O)-;  
each X independently is -O-, -S-, -S(O)-, -S(O)<sub>2</sub>-, -C(O)-, -NR<sup>8</sup>-;  
n is 3 or 4;  
each Y independently is -O-, -S-, -S(O)-, -S(O)<sub>2</sub>-, -C(O)-, -NR<sup>8</sup>-;  
20 m is 2 or 3;  
each Z is -O-C(O)-, -C(O)-O-, -NH-C(O)-, -C(O)-NH-, -O-S(O)<sub>2</sub>-;  
t is 1 or 2;  
R<sup>8</sup> is hydrogen,  $\text{C}_{1-6}$ alkyl,  $\text{C}_{1-6}$ alkylcarbonyl or  $\text{C}_{1-6}$ alkyl-S(O)-;  
R<sup>5</sup> and R<sup>6</sup> each independently are hydrogen, halo, hydroxy,  $\text{C}_{1-6}$ alkyl,  $\text{C}_{1-6}$ alkyloxy,  
25 aryloxy or arylmethoxy;  
R<sup>7</sup> is hydrogen;  
Alk<sup>1</sup> is  $\text{C}_{2-5}$ alkanediyl;  
Alk<sup>2</sup> is  $\text{C}_{2-15}$ alkanediyl;  
Q is a radical of formula
- 30

-27-



wherein

 $R^9$  is hydrogen, cyano, aminocarbonyl or  $C_{1-6}$ alkyl; $R^{10}$  is hydrogen,  $C_{1-6}$ alkyl,  $C_{3-6}$ alkenyl,  $C_{3-6}$ alkynyl,  $C_{3-6}$ cycloalkyl or aryl $C_{1-6}$ alkyl;15  $R^{11}$  is hydrogen or  $C_{1-6}$ alkyl; or $R^{10}$  and  $R^{11}$  taken together may form a bivalent radical of formula  $-(CH_2)_4-$  or  $-(CH_2)_5-$ , or a piperazine which is optionally substituted with  $C_{1-6}$ alkyl; $R^{12}$ ,  $R^{13}$ ,  $R^{14}$ ,  $R^{19}$ ,  $R^{20}$ ,  $R^{21}$ ,  $R^{22}$ ,  $R^{23}$ ,  $R^{24}$ ,  $R^{25}$ ,  $R^{26}$ ,  $R^{27}$ ,  $R^{28}$ ,  $R^{29}$ ,  $R^{36}$ ,  $R^{37}$  and  $R^{38}$  each independently are hydrogen, hydroxy, halo,  $C_{1-6}$ alkyl,  $C_{1-6}$ alkyloxy, aryloxy,  $C_{1-6}$ alkylthio, cyano, amino, mono- or di( $C_{1-6}$ alkyl)amino, mono- or di( $C_{3-6}$ cycloalkyl)-amino, aminocarbonyl,  $C_{1-6}$ alkyloxycarbonylamino,  $C_{1-6}$ alkylaminocarbonylamino, piperidinyl, pyrrolidinyl;20  $R^{15}$ ,  $R^{18}$  and  $R^{35}$  each independently are hydrogen,  $C_{1-6}$ alkyl,  $C_{1-6}$ alkylcarbonyl, or aryl $C_{1-6}$ alkyl;25  $q$  is 1, 2 or 3; $R^{16}$  and  $R^{17}$  are both hydrogen, or taken together with the carbon atom to which they are

connected form C(O);

r is 1, 2 or 3;

R<sup>30</sup> and R<sup>31</sup> are both hydrogen or taken together with the carbon atom to which they are connected form C(O);

5 R<sup>32</sup> is hydrogen, halo or C<sub>1-6</sub>alkyl;

R<sup>33</sup> is hydrogen and R<sup>34</sup> is hydroxy; or R<sup>33</sup> and R<sup>34</sup> taken together may form a bivalent radical of formula (CH<sub>2</sub>)<sub>3</sub> or (CH<sub>2</sub>)<sub>4</sub> which is optionally substituted with C<sub>1-6</sub>alkyl; and aryl is phenyl optionally substituted hydroxy, halo, C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkyloxy.

10 2. A compound as described in claim 1; with the proviso that

(a) N-[2-[2-(2-methoxyphenoxy)ethylamino]ethyl]guanidine; and

(b) the compounds of formula (I) wherein R<sup>3</sup> is methoxy, ethoxy or isopropyl; R<sup>4</sup> is hydrogen; R<sup>5</sup> is hydrogen; R<sup>6</sup> is chloro, fluoro or methyl; R<sup>7</sup> is hydrogen; R<sup>2</sup> is hydrogen or methyl; R<sup>1</sup> is hydrogen; Alk<sup>1</sup> is 1,2-ethanediyl or 1,3-propanediyl; Alk<sup>2</sup> is

15 1,2-ethanediyl or 1,3-propanediyl; Q is a radical of formula (bb), wherein R<sup>12</sup> is hydrogen and R<sup>13</sup> is 4-aminocarbonyl are excluded.

3. A compound as claimed in claim 2, wherein R<sup>3</sup> is C<sub>1-6</sub>alkyl, hydroxy,

20 C<sub>1-6</sub>alkyloxy, aryloxy, arylmethoxy, C<sub>2-6</sub>alkenyl, C<sub>2-6</sub>alkynyl; one of R<sup>4</sup>, R<sup>5</sup> and R<sup>6</sup> is hydrogen and the others each independently are hydrogen, halo, hydroxy, C<sub>1-6</sub>alkyl, or C<sub>1-6</sub>alkyloxy, Q is a radical of formula (aa), (bb), (cc), (dd), (ee) wherein R<sup>38</sup> is hydrogen, (ff), (gg), (hh), (ii), (jj), (kk), (ll).

4. A compound as claimed in claim 2, wherein R<sup>3</sup> and R<sup>4</sup> taken together form a

25 bivalent radical of formula

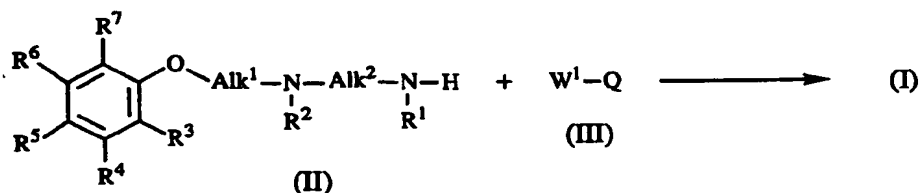
-CH=CH-CH=CH-	(a),	-X-CH=CH-	(f),
-(CH <sub>2</sub> ) <sub>n</sub> -	(b),	-O-(CH <sub>2</sub> ) <sub>t</sub> -Y-	(g),
-(CH <sub>2</sub> ) <sub>m</sub> -X-	(c),	-Y-(CH <sub>2</sub> ) <sub>t</sub> -O-	(h),
-X-(CH <sub>2</sub> ) <sub>m</sub> -	(d),	-(CH <sub>2</sub> ) <sub>t</sub> -Z-	(i),
-CH=CH-X-	(e),	-Z-(CH <sub>2</sub> ) <sub>t</sub> -	(j),

in these bivalent radicals one or two hydrogen atoms may be substituted with C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkylcarbonyl or C<sub>1-6</sub>alkylsulfoxyl; X, Y, Z, m and n are defined as in claim 1, in the bivalent radicals of formula (g) and (h) t is 2, and Q is a radical of formula (aa), (bb), (cc), (dd), (ee) wherein R<sup>38</sup> is hydrogen, (ff), (gg), (hh), (ii), (jj), (kk), (ll).

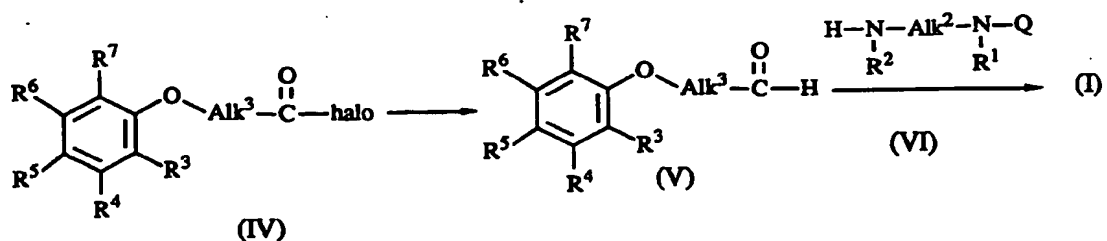
30

5. A compound as claimed in claim 2 wherein R<sup>1</sup> and R<sup>2</sup> are hydrogen.

6. A compound according to claim 2, wherein the compound is  
 N-[2-(2,3-dimethoxyphenoxy)ethyl]-N'-2-pyrimidinyl-1,3-propanediamine;  
 2-[2-[[3-(2-pyrimidinylamino)propyl]amino]ethoxy]phenol; N-[2-(2,3-dimethoxy-  
 phenoxy)ethyl]-N'-(1,4,5,6-tetrahydro-2-pyrimidinyl)-1,3-propanediamine;  
 N-[2-(2-methoxyphenoxy)ethyl]-N'-(1,4,5,6-tetrahydro-2-pyrimidinyl)-1,3-propane  
 diamine; N-[2-(2-ethoxyphenoxy)ethyl]-N'-(1,4,5,6-tetrahydro-2-pyrimidinyl)-  
 1,3-propanediamine; N-[3-(2-methoxyphenoxy)propyl]-N'-(1,4,5,6-tetrahydro-2-  
 pyrimidinyl)-1,3-propanediamine; N-[2-[(2,3-dihydro-1,4-benzodioxin-5-yl)oxy]-  
 ethyl]-N'-2-pyrimidinyl-1,3-propanediamine; N-[2-[(2,3-dihydro-1,4-benzodioxin-5  
 yl)-oxy]ethyl]-N'-(1,4,5,6-tetrahydro-2-pyrimidinyl)-1,3-propanediamine;  
 N-[2-[(2,3-dihydro-1,4-benzodioxin-5-yl)oxy]ethyl]-N'-(1,4,5,6-tetrahydro-2-  
 pyrimidinyl)-1,4-butanediamine; N-[2-(1-naphthalenyloxy)ethyl]-  
 N'-(1,4,5,6-tetrahydro-2-pyrimidinyl)-1,3-propanediamine, a pharmaceutically  
 acceptable acid addition salt thereof or a stereochemically isomeric forms thereof.
7. A pharmaceutical composition comprising a pharmaceutically acceptable carrier and as  
 an active ingredient a therapeutically effective amount of a compound as claimed in  
 claim 2.
8. A process of preparing a composition as claimed in claim 7 characterized in that a  
 therapeutically active amount of a compound as claimed in claim 1 is intimately mixed  
 with a pharmaceutically acceptable carrier.
9. A compound as claimed in claim 2 for use as a medicine.
10. A process of preparing a compound as claimed in claim 2, characterized by  
 a) reacting an intermediate of formula (II), wherein R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, Alk<sup>1</sup>  
 and Alk<sup>2</sup> are as defined in claim 1, with a reagent of formula (III), wherein Q is as  
 defined in claim 1 and W<sup>1</sup> is a reactive leaving group;

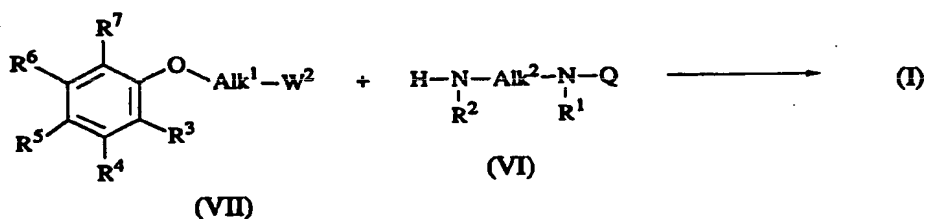


b) reducing an acyl derivative of formula (IV), wherein  $R^3$ ,  $R^4$ ,  $R^5$ ,  $R^6$  and  $R^7$  is as defined in claim 1,  $Alk^3$  is  $C_{1-4}$ alkanediyl, and reacting the resulting aldehyde of formula (V) with an intermediate of formula (VI), wherein  $R^1$  and  $R^2$  are as defined in claim 1.



5

c) *N*-alkylating an amine of formula (VI) with an intermediate of formula (VII), wherein  $R^3$  to  $R^7$  and  $Alk^1$  are as defined in claim 1 and  $W^2$  is a reactive leaving group.



10

and optionally converting the compounds of formula (I) into each other by a functional group transformation reaction; and, if desired, converting a compound of formula (I) into a therapeutically active non-toxic acid addition salt, or conversely, converting an acid addition salt into a free base form with alkali; and/or preparing stereochemically isomeric forms thereof.

15



## INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 94/02702

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07D239/42 C07D319/18 C07D405/12 C07D409/12 C07D239/14  
A61K31/505

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	ARZNEIMITTEL FORSCHUNG. DRUG RESEARCH, vol.25, no.9, 1975, AULENDORF DE pages 1404 - 1408 B.BENKERT 'BEZIEHUNG ZWISCHEN STRUKTUR UND NORADRENALIN-ENTSPICHERNDER WIRKUNG VON GUANIDIN-UND AMIDINDERIVATEN.' cited in the application see page 1404 - page 1407 ---	1,7-9
A	US,A,4 593 039 (JOHN J. BALDWIN ET AL.) 3 June 1986 see column 1 - column 19 ---	1-9
A	EP,A,0 511 072 (SYNTHELABO) 28 October 1992 cited in the application see page 1 - page 9 ---	1-9
-/--		

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

## \* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

24 November 1994

Date of mailing of the international search report

06.12.94

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
Fax: (+31-70) 340-3016

Authorized officer

Francois, J

# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/EP 94/02702

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>EP,A,0 042 593 (BOEHRINGER) 30 December 1981  see page 1 - page 9; claims; example 3  -----</p>	1-9

# INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

PCT/EP 94/02702

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US-A-4593039	03-06-86	US-A- 4810725	07-03-89
EP-A-0511072	28-10-92	FR-A- 2675799	30-10-92
		AU-A- 1507992	29-10-92
		CN-A- 1066066	11-11-92
		JP-A- 5117243	14-05-93
		US-A- 5229392	20-07-93
EP-A-0042593	30-12-81	DE-A- 3023369	14-01-82
		AT-T- 11535	15-02-85
		AU-B- 540425	15-11-84
		AU-A- 7187981	07-01-82
		CA-A- 1184188	19-03-85
		EP-A, B 0042592	30-12-81
		JP-B- 2003778	24-01-90
		JP-A- 57045144	13-03-82
		JP-A- 57032269	20-02-82
		SU-A- 1287748	30-01-87
		SU-A- 1272976	23-11-86
		US-A- 4438128	20-03-84
		US-A- 4507488	26-03-85
		US-A- 4608383	26-08-86